

CANTHARIS- cantharis gel
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

ACTIVE INGREDIENT

CANTHARIS GEL 3X HPUS

USES

Minor burns and scalds

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE & ADMINISTRATION SECTION

Apply a thin layer of Gel to the affected area, repeat 3 times a day or as needed.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are pregnant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.
Close the cap tightly after use.

INACTIVE INGREDIENTS

ALOE VERA GEL

STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com

Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758

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CANTHARIS GEL



Manufactured according to HPUS
FDA Est. # 30052969310 | India ML 1/DH/14 | Rxhomeo.com
Ingredients: Cantharis 3X 10% in Aloe Vera Gel Base

USES: Minor burns and scalds

Distributed in the US by Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX, 78758 Manufactured by Rxhomeo Private Limited "Indrashanah" 4-1-424 to 426, Bank Street, Abids, Hyderabad #500001 India

NDC 15631-2311-0
B.No 23110018C1
MFD XXXX EXP XXXX
Contents 50 Grams



EXTERNAL APPLICATION ONLY

CANTHARIS

cantharis gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15631-2311
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	3 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15631-2311-0	1 in 1 CONTAINER	03/30/2016	
1		50 g in 1 CONTAINER; Type 0: Not a Combination Product		
2	NDC:15631-2311-1	1 in 1 CONTAINER	03/30/2016	
2		100 g in 1 CONTAINER; Type 0: Not a Combination Product		
3	NDC:15631-2311-2	1 in 1 CONTAINER	03/30/2016	
3		200 g in 1 CONTAINER; Type 0: Not a Combination Product		
4	NDC:15631-2311-3	1 in 1 CONTAINER	03/30/2016	
4		500 g in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/30/2016	

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-2311) , label(15631-2311)

Revised: 3/2016

Rxhomeo Private Limited d.b.a. Rxhomeo, Inc